

02/12/99
JC525 U.S. PTO

Express Mail No. EM036758282US
Docket No. A 12222 US (C36368/108310)

Assistant Commissioner for Patents
Box Patent Application
Washington, D.C. 20231

Sir:

Transmitted herewith for filing is the utility patent application of Inventors:

DENISE ANDERSON and GEORG FRÄTER

For: ARYL-ACRYLIC ACID ESTERS

1. The application has 35 pages (including claim pages, attached declaration and power of attorney, and abstract).
2. ____ sheet(s) of drawing is (are) enclosed. The drawings are:
 - a. formal; or
 - b. informal; formal drawings will be submitted in due course.
3. The declaration and power of attorney
 - a. has been executed by all the inventors; or
 - b. has not been executed by all the inventors. A signed declaration and power of attorney will be submitted in due course.
4. An associate power of attorney is enclosed.
5. An assignment of the invention to Givaudan Roure (International) S.A. and a Recordation Form Cover Sheet are enclosed. Please record the Assignment and return it to the undersigned. A duplicate copy of this paper is enclosed.
 - a. A check for \$40.00 to cover the recording fee is enclosed. See paragraph 10.c., below.
 - b. Please charge the recording fee to our Deposit Account No. 02-4467. A duplicate copy of this paper is enclosed.
6. Priority is hereby claimed under 35 USC § 119 based on Appln. No. 98810114.3, filed February 13, 1998, in Europe.
 - a. A certified copy of each of the priority documents is enclosed.
 - b. The certified priority document(s) will follow.

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7. A verified statement to establish small entity status under 37 CFR Section 1.9 and 37 CFR Section 1.27 is enclosed.

8. The filing fee is calculated below.

For	<u>Col. 1</u> No. Filed		<u>Col. 2</u> No. Extra	<u>Small Entity</u> Rate Fee	or	<u>Other Than A</u> <u>Small Entity</u> Rate Fee
Basic Fee:				\$ 385	or	\$ 760
Total Claims:	18-20=	x	0	x11=	or	x22=0
Indep. Claims	2-3=	x	0	x41=	or	x82=0
[] Multiple Dependent Claims Presented				+ 130 =	or	+ 260 = 0
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9. A Preliminary Amendment is enclosed.

a. No additional fee is due.

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10. The Commissioner is hereby authorized to charge payment of the following fees associated with this application or with recording any Assignment concerning it, or to credit any overpayment, to Deposit Account No. 02-4467, unless otherwise paid by check.

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- d. [x] Any additional Assignment recording fees under 37 CFR Section 1.21(h).

Note: See 37 CFR § 1.311(b) regarding authorization to pay the issue fee from deposit account.

- 11. [] An Information Disclosure Statement is enclosed.

Respectfully submitted,

By: 
Warren K. MacRae
Registration No. 37,876
BRYAN CAVE LLP
245 Park Avenue
New York, New York 10167-0034
Phone: (212) 692-1800
Fax: (212) 692-1900

ARYL-ACRYLIC ACID ESTERS

5

FIELD OF INVENTION

The present invention relates to 3-(2-substituted aryl)-acrylic acid esters, especially 3-(2-hydroxyaryl)-acrylic acid esters and 3-(2-aminoaryl)-acrylic acid esters. These 10 compounds are useful as precursors for organoleptic compounds, such as for example, flavors, fragrances and masking agents, antimicrobial compounds and fluorescent whitening agents.

BACKGROUND OF THE INVENTION

15

A principal strategy currently employed for imparting odors, such as fragrance, to consumer products is to admix the fragrance directly into the product. There are, however, several drawbacks to this strategy. For example, the fragrance 20 material can be too volatile and/or too soluble, resulting in fragrance loss during manufacturing, storage, and use. In addition, many fragrance materials are also unstable over time which, again, results in loss during storage.

25

In many consumer products it is desirable for the fragrance to be released slowly over time. Microencapsulation and inclusion complexes with cyclodextrins have been used to help decrease volatility, improve stability and provide slow-release properties. These methods are, for a number of 30 reasons, often not successful and/or too expensive to be commercially viable, as for example, in the case of cyclodextrins.

Fragrance precursors for scenting fabrics being washed in the presence of a lipase-containing detergent are described in WO 95/04809. The fragrance precursors contained in the detergent and/or in the softener are cleaved by the lipase and 5 a single active compound. In such a situation, an odoriferous alcohol, aldehyde or ketone is produced. In this system, a prolonged scenting effect on the fabric is obtained. However, the need for a lipase-containing detergent is of limited commercial significance. In many parts of the world, 10 detergents do not contain lipase. Other consumers prefer to use so-called 'nonbio' detergents.

Fluorescent whitening agents or brighteners have been added to laundry detergents since the 1950s to help maintain 15 the original brightness of white clothing.

Accordingly, one object of the present invention is to provide new precursors for compounds with different activities. Through these precursors, different activities 20 are imparted to a product by the addition of just one compound of the present invention.

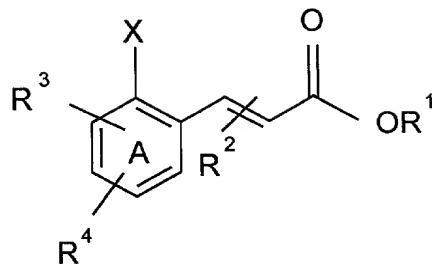
Another object of the invention is to provide new precursor compounds which are stable under transport and 25 storage conditions.

A further object of the present invention is to provide new precursors that supply different active compounds simultaneously or successively.

30

SUMMARY OF THE INVENTION

The present invention provides compounds of the Formula I



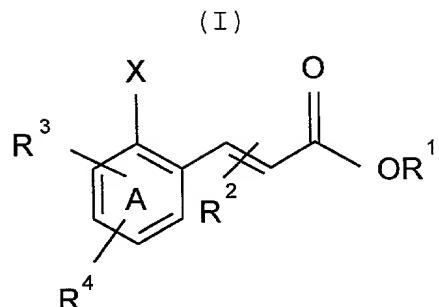
wherein A is a benzene or a naphthalene ring; R¹ is, or is selected from saturated, unsaturated, straight, branched, 5 alicyclic and/or aromatic C₁₀-C₃₀ hydrocarbon residues which can contain heteroatoms and can be substituted by an ionic substituent; R² in the 2- or 3-position is, or is selected from hydrogen, a straight or branched C₁-C₆ residue, an 10 optionally substituted aromatic and/or optionally substituted heterocyclic residue; R³ and R⁴ are, or selected from hydrogen, a straight or branched C₁-C₆ alkyl, a C₁-C₆ alkoxy residue, a substituted or condensed heterocyclic residue, -OH, -NO₂, -NH₂, -N(C₁-C₆ alkyl)₂, -N(hydroxyalkyl)₂, -NHCO₂CH₃ and/or -NH(heterocycle). 15

In the present invention, R², R³ and R⁴ are the same or different. X is, or is selected from -OH and/or NHR⁶, wherein R⁶ is, or is selected from hydrogen, a saturated or unsaturated, straight or branched C₁-C₂₀ hydrocarbon and/or an 20 optionally substituted aromatic or heterocyclic residue; and the acrylic double bond is of the E configuration.

The present invention also provides a method for preparing organoleptic, antimicrobial and fluorescent 25 whitening compositions by incorporating into these precursor compounds of Formula I described above.

DETAILED DESCRIPTION OF THE INVENTION

The present invention provides novel 3-(2-substituted aryl)-acrylic acid esters of the Formula I set forth below:



5

wherein

A is a benzene or naphthalene ring;

10 R¹ is a saturated or unsaturated, straight, branched, alicyclic or aromatic C₁₀-C₃₀ hydrocarbon residue which can contain heteroatoms and can be substituted by an ionic substituent;

15 R² in the 2- or 3-position is hydrogen, a straight or branched C₁-C₆ residue, an optionally substituted aromatic or an optionally substituted heterocyclic residue;

20 R³ and R⁴ are hydrogen, a straight or branched C₁-C₆ alkyl or C₁-C₆ alkoxy residue, a substituted or condensed heterocyclic residue, -OH, -NO₂, -NH₂, -N(C₁-C₆ alkyl)₂, -N(hydroxyalkyl)₂, -NHCO₂CH₃ or -NH(heterocycle);

25 R², R³ and R⁴ may be the same or different;

X is -OH or NHR⁶, wherein R⁶ is hydrogen a saturated or unsaturated, straight or branched C₁-C₂₀ hydrocarbon or an optionally substituted aromatic or heterocyclic residue; and

the acrylic double bond is of the E configuration.

In the present invention, Formula I includes all possible
5 enantiomers, diastereomers and all mixtures thereof.

In Formula I, R^1 is preferably a saturated or unsaturated, straight or branched $C_{10}-C_{30}$ hydrocarbon residue containing one or more O and/or N atoms and/or $C(O)$ groups
10 and/or alkoxy groups or substituted by an ionic substituent of the formula $N(R^5)_3^+$, in which R^5 is the residue of a fatty acid or an alkyl group with 1 to 30 carbon atoms are preferred.

In Formula I, R^2 is preferably a heterocyclic residue of
15 the formula



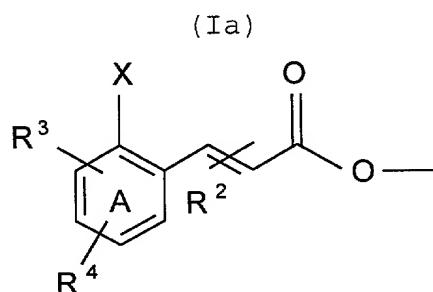
In Formula I, R^3 and/or R^4 are preferably hydrogen, $-N(C_1-C_6$ alkyl) $_2$, $-NH_2$, a five membered heterocyclic residue
20 optionally containing N and/or O atoms, substituted by C_1-C_6 aliphatic and/or aromatic substituents.

In one embodiment of the present invention, R^1 is
25 preferably the residue of an olfactory alcohol of the formula R^1OH , of the enol form of an olfactory aldehyde of the formula R^1HO or of an olfactory ketone of formula R^1O . Alternatively,
R¹ is an optionally substituted alkyl, alkenyl or arylalkyl
30 residue carrying an 1-alkoxy, 1-aryloxy, or 1-arylalkoxy residue.

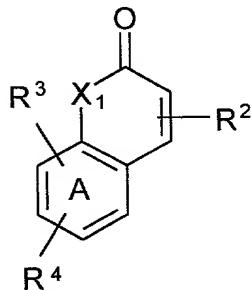
In the present invention, compounds within the scope of Formula I are mostly or nearly odorless at room temperature, atmospheric conditions and about 20 to 100 % relative humidity. Under activating conditions, however, compounds of 5 Formula I are cleaved to form active compounds. For example, the residue of Formula Ia when cleaved forms a coumarin of Formula II.

The structures of Formulae Ia and II are set forth below:

10



(II)



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wherein X^1 is O or NR^6 ; R^2 , R^3 , R^4 are defined as set forth above.

20

In the present invention, the coumarins of Formula II have organoleptic, antimicrobial and/or optical brightening activity. Preferably, coumarins of the present invention have olfactory properties.

In the present invention, when R¹ is the residue of an olfactory alcohol or the enol form of an olfactory aldehyde or ketone, upon cleavage two active compounds with the above properties can be obtained. Thus, the compounds of Formula I 5 permit the development of useful consumer products with enhanced organoleptic, antimicrobial and/or optical brightening properties. The organoleptic coumarins, alcohols, aldehydes and ketones obtained from the present compounds are useful, for example, in preparing fragrances, flavors, masking 10 agents and antimicrobial agents.

To form active residues from Formula I and Ia, these compounds are subject to activating conditions that lead to the cleavage of thereof. In the present invention, actinic 15 radiation, such as, for example, UV light, including sunlight can be used to activate the compounds of Formula I. Elevated temperatures also function to activate the compounds of Formula I and Ia. For purposes of the present invention, any activating condition may be used, so long as the desired 20 active ingredients are formed by cleavage of compounds of Formula I and Ia to form the desired residue with the proper activity.

Another embodiment of the present invention is a method 25 for preparing organoleptic, antimicrobial and fluorescent whitening compositions wherein the compounds of Formula I are precursors for these compounds. For purposes of the present invention, organoleptic precursors include, for example, precursors compatible in flavors, fragrances and odor masking 30 agents.

In the present invention, the esters of formula I can act as fragrance precursors in laundry products. They can also act as precursors for odor masking agents in the same products as

the fragrance precursors. They also can act as precursors for antimicrobial agents. In addition, they can also act as precursors for fluorescent whitening agents. The fragrance precursors and the precursors for odor masking agents as well 5 as the flavor precursors of the invention may be used individually in an amount effective to enhance or to mask the characteristic odor or flavor of a material. More commonly, however, the compounds are mixed with other fragrance or flavor components in an amount sufficient to provide the 10 desired odor or flavor characteristics.

Thus, as used herein, an "effective" amount is intended to mean a sufficient quantity of the precursor to obtain the desired effect, i.e., fragrance, odor masking and/or 15 flavoring. Reference is made to the Examples set forth below and to the skill of the art for determining an effective amount of the present inventive compounds in any given situation.

20 The brightener precursors may also be employed individually in an effective amount where appropriate, and/or mixed with one or more other brightener or colorant substance.

Due to the *in situ* generation of the active compounds of 25 the present invention, the desired effect is prolonged and the substantivity on different substrates is enhanced. If two active compounds are provided by one ester of the Formula I, they can be generated, depending on the precursor and/or the activating conditions, simultaneously or successively. 30 Further, the precursors of the present invention provide slow release of the active compounds.

Non-limiting examples of alcohols R^1OH generated upon cleavage of compounds of Formula I and constituting the

residue R¹, include amyl alcohol, hexyl alcohol*, 2-hexyl alcohol*, heptyl alcohol*, octyl alcohol*, nonyl alcohol*, decyl alcohol* undecyl alcohol*, lauryl alcohol*, myristic alcohol, 3-methyl-but-2-en-1-ol*, 3-methyl-1-pentanol, cis-3-5 hexenol*, cis-4-hexenol*, 3,5,5-trimethyl-hexanol, 3,4,5,6,6-pentamethylheptan-2-ol*, citronellol*, geraniol*, oct-1-en-3-ol, 2,5,7-trimethyl-octan-3-ol, 2-cis-3,7-dimethyl-2,6-octadien-1-ol, 6-ethyl-3-methyl-5-octen-1-ol*, 3,7-dimethyl-10 oct-3,6-dienol*, 3,7-dimethyloctanol*, 7-methoxy-3,7-dimethyl-octan-2-ol*, cis-6-nonanol*, 5-ethyl-2-nananol, 6,8-dimethyl-2-nananol*, 2,2,8-trimethyl-7(8)-nonene-3-ol, nona-2,6-dien-1-ol, 4-methyl-3-decen-5-ol*, dec-9-en-1-ol, benzylalcohol, 2-methyl-undecanol, 10-undecen-1-ol, 1-phenyl-ethanol*, 2-phenyl-ethanol*, 2-methyl-3-phenyl-3-propenol, 2-phenyl-15 propanol*, 3-phenyl-propanol*, 4-phenyl-2-butanol, 2-methyl-5-phenyl-pentanol*, 2-methyl-4-phenyl-pentanol*, 3-methyl-5-phenyl-pentanol*, 2-(2-methylphenyl)-ethanol*, 4-(1-methylethyl)-benzene-methanol, 4-(4-hydroxyphenyl)-butan-2-one*, 2-phenoxy-ethanol*, 4-(1-methylethyl)-2-hydroxy-1-methyl-20 benzene, 2-methoxy-4-methyl-phenol, 4-methyl-phenol, anisic alcohol*, p-tolyl alcohol*, cinnamic alcohol*, vanillin*, ethyl vanillin*, eugenol*, isoeugenol*, thymol, anethol*, decahydro-2-naphthalenol, borneol*, cedrenol*, farnesol*, fenchyl alcohol*, menthol*, 3,7,11-trimethyl-2,6,10-dodecatrien-1-ol, alpha ionol*, tetrahydro ionol*, 2-(1,1-dimethylethyl)cyclohexanol*, 3-(1,1-dimethylethyl)cyclohexanol*, 4-(1,1-dimethylethyl)cyclohexanol*, 4-isopropyl-cyclohexanol, 6,6-dimethyl-bicyclo[3.3.1]hept-2-ene-2-ethanol, 6,6-dimethyl-30 bicyclo[3.1.1]hept-2-ene-methanol*, p-menth-8-en-3-ol*, 3,3,5-trimethyl-cyclohexanol, 2,4,6-trimethyl-3-cyclohexenyl-methanol*, 4-(1-methylethyl)-cyclohexyl-methanol*, 4-(1,1-dimethylethyl)-cyclohexanol, 2-(1,1-dimethylethyl)-cyclohexanol, 2,2,6-trimethyl-alpha-propyl-cyclohexane

propanol*, 5-(2,2,3-trimethyl-3-cyclopentenyl)-3-methylpentan-2-ol*, 3-methyl-5-(2,2,3-trimethylcyclopentyl-3-enyl)pent-4-en-2-ol*, 2-ethyl-4(2,2,3-trimethylcyclopentyl-3-enyl)but-2-en-1-ol*, 4-(5,5,6-trimethylbicyclo[2.2.1]hept-2-yl)-5-cyclohexanol*, 2-(2-methylpropyl)-4-hydroxy-4-methyl-tetrahydropyran*, 2-cyclohexyl-propanol*, 2-(1,1-dimethylethyl)-4-methyl-cyclohexanol*, 1-(2-tert-butyl-cyclohexyloxy)-2-butanol*, 1-(4-isopropyl-cyclohexyl)-ethanol*, 2,6-dimethyl-oct-7-en-2-ol**, 2,6-dimethyl-heptan-2-ol** and 3,7-dimethyl-octa-1,6-dien-3-ol**.

In the list set forth above, the alcohols identified with an asterisk are preferred embodiments, and the alcohols identified with two asterisks are more preferred.

Non-limiting examples of aldehydes R^1HO generated upon cleavage a compound of Formula I and constituting the residue R^1 thereof include, for example, 2,6,10-trimethylundec-9-enal*, 1,2,3,4,5,6,7,8,-octahydro-8,8-dimethyl-2-naphthalenecarboxaldehyde tridecanal, 2-[4-(1-methylethyl)phenyl]-ethanal, 2,4-dimethyl-cyclohex-3-ene-1-carbox-aldehyde*, 4-carboxaldehyde-1,3,5-trimethyl-cyclohex-1-ene*, 1-carboxaldehyde-2,4-dimethyl-cyclohex-3-ene*, 1-carboxaldehyde-4-(4-hydroxy-4-methylpentyl)-cyclohex-3-ene*, 3,5,5-trimethyl-hexanal, heptanal*, 2,6-dimethyl-hept-5-enal*, decanal**, dec-9-enal, dec-4-enal, 2-methyldecanal*, undec-10-enal**, undecanal*, dodecanal**, 2-methyl-undecanal**, tridecanal, octanal**, nonanal*, 3,5,5-trimethylhexanal, undec-9-enal**, 2-phenyl-propanal*, 4-methyl-phenyl-acetaldehyde*, 3,7-dimethyl-octanal*, dihydrofarnesal**, 7-hydroxy-3,7-dimethyl-octanal*, 2,6-dimethyl-oct-5-enal, 2-[4-(1-methylethyl)phenyl]-ethanal*, 3-(3-isopropyl-phenyl)-butanal**, 2-(3,7-dimethyoct-6-enoxy)-ethanal, 1-carboxaldehyde-4-(4-methyl-3-pentenyl)-cyclohex-3-

ene*, 2,3,5,5,-tetramethyl-hexanal, longifolic aldehyde, 2-methyl-4-(2,6,6-trimethylcyclohex-2-en-1-yl)-butanal*, 2-methyl-3-(4-tert-butylphenyl)-propanal**, 4-(1,1-dimethyl-ethyl)-benzene-propanal*, 2-[4-(1-methyl-ethyl)-phenyl]-5-propanal, alpha-methyl-1,3-benzodioxole-5-propanal*, 3,7-dimethyl-oct-6-enal*, 2-methyl-3-(4-isopropylphenyl)-propionaldehyde*, 4-(4-hydroxy-4-methyl-pentyl)-cyclohex-3-en-1-carboxaldehyde**, alpha-methyl-1,3-benzodioxole-5-propanal*, 1-carboxaldehyde-4-(1,1-dimethylethyl)-cyclohexane, 4-(octahydro-4,7-methano-5H-inden-5-ylidene)-butanal and [(3,7-dimethyl-6-octenyl)-oxy]-acetaldehyde**.

In the list set forth above, the aldehydes identified with an asterisk are preferred embodiments, and the alcohols 15 identified with two asterisks are more preferred.

Non-limiting examples of ketones R^1O generated upon cleavage a compound of Formula I and constituting the residue R^1 thereof include, for example, 2-heptyl-cyclopentanone, 20 2,2,6,10-tetramethyltricyclo-[5.4.0.0(6,10)]-undecan-4-one benzylacetone*, carvone*, 1,2,3,5,6,7-hexahydro-1,1,2,3,3,-pentamethyl-4H-inden-4-one*, methyl heptenone*, dimethyl octenone*, 2-(butan-2-yl)-cyclohexanone*, 2-hexyl-cyclopent-2-en-1-one*, 2-(1-methylethyl)-5-methyl-cyclohexanone*, 2-(2-methylethyl)-5-methyl-cyclohexanone*, 3-methyl-25 cyclopentadecanone, 4-tert-pentyl-cyclohexanone*, 3-oxo-2-pentyl-cyclopentane-acetic acid methyl ester**, 1-(1,2,3,4,5,6,7,8-octahydro-2,3,8,8-tetramethyl-2-naphthalenyl)-ethanone* and 3-methyl-5-propyl-cyclohex-2-en-1-one*.

In the list set forth above, the ketones identified with an asterisk are preferred embodiments, and the alcohols identified with two asterisks are more preferred.

Non-limiting examples of fluorescent whitening coumarins generated upon cleavage a compound of Formula I and constituting the residue R¹ thereof include, for example, 7-
5 (3-methyl-1H-pyrazol-1-yl)-3-phenyl-2H-1-benzopyran-2-one, 7-
(4-methyl-5-phenyl-2H-1,2,3-triazol-2-yl)-3-phenyl-2H-1-
benzopyran-2-one, 7-(2H-naphtho[1,2-d]triazol-2-yl)-3-phenyl-
2H-1-benzo-pyran-2-one, 3-(1H-pyrazol-1-yl)-7-(2H-1,2,3-
10 triazol-2-yl)-2H-1-benzo-pyran-2-one, 7-(dimethylamino)-1-
methyl-3-phenyl-2(1H)-quinolinone, 7-(diethylamino)-1-ethyl-3-
phenyl-2(1H)-quinolinone, 7-amino-4-methyl-2H-1-benzopyran-2-
one, 7-(dimethylamino)-4-methyl-2H-1-benzopyran-2-one, 7-
(diethylamino)-4-methyl-2H-1-benzopyran-2-one, 7-hydroxy-4-
methyl-2H-1-benzopyran-2-one and 6,7-dihydroxy-2H-1-
15 benzopyran-2-one.

Non-limiting examples of coumarins of Formula II having olfactory properties generated upon cleavage and constituting the respective residue in the compounds of Formula I, include
20 2H-1-benzopyran, 3-methyl-benzopyran-2-one, 8-(1,1-dimethylethyl)-6-methyl-benzopyrone, 4-methyl-7-ethoxy-
coumarin and 6-methyl-2H-1-benzopyran.

The active coumarins, alcohols, aldehydes and ketones
25 which are generated as a result of the desired cleavage of the acrylic acid esters of Formula I by UV-light and/or by elevated temperatures is exemplary only and is not intended to limit the scope of the present invention in any way.

30 The compounds of Formula I, preferably, may be employed in preparing sustained release odorants and flavors. The compounds of Formula I also are useful as sustained agents which may be employed to mask or attenuate undesirable odors or to provide additional odors not initially present in

consumer products including for example, laundry detergents, fabric softeners, fabric softener sheets, toiletries and cosmetics such as sunscreens.

5 Further applications of the compounds of Formula I include sustained brighteners and antimicrobial agents in the same products. The brighteners are especially useful for wool, rayon and polyamides. These compounds are also useful for flavoring and aromatizing tobacco products, such as for
10 example, cigarettes.

The amount of the compounds of the present invention required to produce the desired, overall effect varies depending upon the particular compounds of Formula I chosen, 15 the product in which it will be used, and the particular effect desired. The examples teach what amounts are suitable for any desired purpose.

For example, depending upon the selection and 20 concentration of the compound chosen, when a compound of Formula I is added either singly or as a mixture to, for example, a laundry product composition at levels ranging from about 0.001 to about 10 % by weight, a coumarin and if desired an odoriferous alcohol, aldehyde or ketone in an 25 organoleptically effective amount is released when the product is used. These newly formed odorant(s) serve to enhance the odor of the fragrance. Depending on the compound of Formula I, an antimicrobial agent and/or a brightener can be released.

30 Depending upon the selection and concentration, addition of the Formula I compounds, either singly or as a mixture, to cigarette tobacco at levels ranging from about 5 ppm to about 50,000 ppm tends to enhance the smoking flavor and/or mask undesirable smoking odors. An important property of these

Formula I compounds is that the flavorant or odorant is covalently bound as a non-volatile compound and the flavorant or odorant is released only when the tobacco product is ignited and burns.

5

Addition of the compounds of Formula I, either separately, or as a mixture, at levels suitably ranging from about 5 ppm to about 50,000 ppm by weight onto the media enclosing the tobacco, serves to incorporate the 10 odorant/flavorant in the side-stream smoke of the tobacco. Air borne flavorants and/or odorants are thus introduced. This newly formed odorant or flavorant enhances and masks the smoking odors, depending upon selection and use levels of the compounds of Formula I.

15

As is evident from the above compilation of alcohols, aldehydes, ketones and coumarins, a broad range of odorants or flavors or mixtures can be generated from precursors of the invention. While manufacturing compositions, the precursors of 20 the invention may be used according to methods known to the perfumer, such as e.g. from W.A. Poucher, Perfumes, Cosmetics, Soaps, 2, 7th Edition, Chapman and Hall, London 1974 which is hereby incorporated by reference. The fluorescent whitening agents may be added in similar manner.

25

The following examples are provided to further illustrate methods of preparation of the compounds of the present invention, as well as certain physical properties and uses thereof. These examples are illustrative only and are not 30 intended to limit the scope of the invention in any way.

EXAMPLES

Example 1

(E)-3-(2-Hydroxy-phenyl)-2-methyl-acrylic acid ethyl ester

To a solution of 75.0 g (carbethoxyethylidene)triphenyl phosphorane in 350 ml of toluene, 23.2 g of salicylaldehyde 5 was dropped in at 20°C while cooling in an ice-bath. After stirring at room temperature for 90 min., the reaction mixture was diluted with toluene and washed to neutrality with water. The organic phase was dried, filtered and evaporated to dryness. The resulting yellow oil was purified by 10 chromatography to yield 35.5 g of a colourless solid.

NMR (CDCl₃) δ 7.82 (s, 1H), 7.30-6.85 (m, 4H), 6.6 (s, OH), 4.35-4.17 (q, 2H), 2.04 (s, 3H), 1.40-1.28 (t, 3H)

15

Example 2

(E)-3-(2-Hydroxy-phenyl)-acrylic acid methyl ester

According to the procedure of Example 1, (E)-3-(2- 20 hydroxy-phenyl)-2-methyl-acrylic acid methyl ester was prepared from methyl(triphenyl-phosphoranylidene)acetate and salicylaldehyde.

25

Example 3

(E)-3-(2-Hydroxy-phenyl)-acrylic acid ethyl ester

According to the procedure of Example 1, (E)-3-(2- 30 hydroxy-phenyl)-2-methyl-acrylic acid methyl ester was prepared from ethyl(triphenyl-phosphoranylidene)acetate and salicylaldehyde.

Example 4

(E)-3-(2-Hydroxy-phenyl)-2-methyl-acrylic acid

To a solution of 35.5 g of (E)-3-(2-hydroxy-phenyl)-2-methyl-acrylic acid ethyl ester in 600 ml of ethanol, a
5 solution of 10.66 g of potassium hydroxide in 500 ml of water was dropped in. After refluxing for 5 hours, another 5.0 g of potassium hydroxide was put in and the mixture was refluxed for another 19 hours. Then the reaction mixture was cooled down, diluted with ether and washed to pH 4 with HCl 2N and
10 water. The organic phase was dried, filtered and evaporated to dryness. The resulting colourless crystals were not further purified.

NMR (CDCl₃) δ 7.90 (s, 1H), 7.31-6.83 (m, 4H), 2.07 (s, 3H)

15

Example 5

(E)-3-(2-Hydroxy-phenyl)-2-methyl-acrylic acid 3,7
dimethyl-oct-6-enyl ester

20 A solution of 6.0 g of (E)-3-(2-hydroxy-phenyl)-2-methyl-acrylic acid, 5.3 g of citronellol and 1 g of p-toluenesulfonic acid in 150 ml of cyclohexane was refluxed for 6.5 hours using a water separator. Then the reaction mixture was cooled down, diluted with hexane and washed to neutrality with
25 saturated sodium bicarbonate and water. The organic phase was dried, filtered and evaporated to dryness. The resulting yellow oil was purified by chromatography to yield 6.45 g of a colourless oil.

30 NMR (CDCl₃) δ 7.80 (s, 1H), 7.38-6.83 (m, 4H), 6.4 (s, OH), 5.09 (t, 1H) 4.25 (t, 2H), 2.10-1.90 (m, 5H), 1.88-1.08 (m, 11H), 0.93 (d, 3H)

Example 6

(E)-3-(2-Hydroxy-phenyl)-2-methyl-acrylic acid phenethyl ester

According to the same procedure of Example 3, (E)-3-(2-hydroxy-phenyl)-2-methyl-acrylic acid phenethyl ester was prepared from 3-(2-hydroxy-phenyl)-2-methyl-acrylic acid, phenyl ethyl alcohol and p-toluenesulfonic acid.

Example 7

10 3-(4-Diethylamino-2-hydroxy-phenyl)-but-2-enoic acid ethyl ester

To a suspension of 5.27 g ethoxycarbonylmethylene-triphenylphosphorane in 10 ml of toluene, a solution of 2.02 g 15 1-(4-diethylamino-2-hydroxy-phenyl)-ethanone (DE 28 44 606) was dropped in at room temperature. Then the reaction mixture was heated to reflux. After refluxing for 31 hours the mixture was cooled down and evaporated to dryness. The resulting dark oil was purified by chromatography to yield a colourless oil.

20 NMR (CDCl₃) δ 7.22-7.09 (m, 1H), 6.57-6.42 (m, 1H),
6.32-6.20 (m, 2H), 5.01 (s, 1H), 4.18-4.01 (q, 2H), 3.42-3.24 (q, 4H), 2.49 (s, 3H), 1.31-1.08 (m, 9H).

25 The compounds of Examples 1-4 yield upon cleavage organoleptic coumarins, Examples 5 and 6 organoleptic coumarins and organoleptic alcohols and Example 7 a brightener coumarin.

30 Example 8
(E)-3-(2-Hydroxy-5-methyl-phenyl)-acrylic acid ethyl ester

A solution of 193.7 g 3-[5-methyl-2-(tetrahydro-pyran-2-yloxy)-phenyl]-acrylic acid ethyl ester (Bunce, R., Moore, J., Org. Prep. Proc., 29(3), 293 (1997)) and 2 g p-toluenesulfonic acid in 2.5 l of ethanol was stirred at room temperature for 5 24 hours. Then the reaction mixture was concentrated and the residue was diluted with ether, washed with saturated sodium bicarbonate and brine, dried and evaporated to dryness. The resulting yellow solid was recrystallized to yield 97.7 g of colourless crystals.

10

NMR (CDCl₃) δ 8.10-7.93 (d, 1H), 7.27 (s, 1H), 7.10-6.98 (d, 1H), 6.82-6.70 (d, 1H), 6.69-6.55 (d, 1H), 4.38-4.20 (q, 2H), 2.27 (s, 3H), 1.42-1.28 (t, 3H) ppm.

15

(E)-3-(2-Hydroxy-phenyl)-acrylic acid

To a solution of 100.0 g 3-(2-hydroxy-phenyl)-acrylic acid ethyl ester in 500 ml of ethanol, a solution of 50.9 g potassium hydroxide in 500 ml of water was dropped in at room 20 temperature. After stirring at reflux for 28 hours, the reaction mixture was concentrated. The residue was diluted with 500 ml HCl 2N and extracted with ether. The organic phase was washed with 2N HCl and water, dried and evaporated to 25 dryness. The resulting solid was recrystallized to yield 44.3 g of colourless crystals.

30

NMR (DMSO) δ 12.2 (br s, 1H), 10.2 (br s, 1H), 7.95-7.75 (d, 1H), 7.65-7.50 (d, 1H), 7.32-7.15 (m, 1H), 6.99-6.76 (m, 2H), 6.62-6.45 (d, 1H) ppm.

(E)-3-(2-Hydroxy-phenyl)-acrylic acid dec-9-enyl ester

A mixture of 5.0 g 3-(2-hydroxy-phenyl)-acrylic acid ethyl ester, 6.1 g dec-9-en-1-ol and 1.0 g tetraisopropyl-ortho-titanate was heated to 150°C removing the ethanol formed. After stirring for 2.5 hours at this temperature, the 5 reaction mixture was cooled, diluted with ether and washed with brine. The organic phase was dried and evaporated to dryness. The resulting oil was Kugelrohr-distilled, crystallized and recrystallized to yield 1.69 g of colourless crystals.

10

NMR (CDCl₃) δ 8.11-7.96 (d, 1H), 7.55-7.40 (d, 1H),
7.32-7.15 (m, 1H), 6.99-6.72 (m, 2H),
6.71-6.57 (d, 1H), 5.93-5.69 (m, 1H),
5.07-4.88 (m, 2H), 4.31-4.15 (t, 2H),
2.12-1.93 (m, 2H), 1.85-1.55 (m, 2H),
1.54-1.15 (m, 10H) ppm.

(E)-3-(2-Hydroxy-phenyl)-acrylic acid 2-ethyl-4-(2,2,3-trimethyl-cyclopent-3-enyl)-but-2-enyl ester

20

According to the same procedure, 3-(2-hydroxy-phenyl)-acrylic acid 2-ethyl-4-(2,2,3-trimethyl-cyclopent-3-enyl)-but-2-enyl ester was prepared from 3-(2-hydroxy-phenyl)-acrylic acid ethyl ester, 2-ethyl-4(2,2,3-trimethyl cyclopentyl-3-en-25 1-yl)-but-2-en-1-ol and tetraisopropyl-o-titanate.

(E)-3-(2-Hydroxy-5-methyl-phenyl)-acrylic acid 2-ethyl-4-(2,2,3-trimethyl-cyclopent-3-enyl)-but-2-enyl ester

30

According to the same procedure, 3-(2-hydroxy-5-methyl-phenyl)-acrylic acid 2-ethyl-4-(2,2,3-trimethyl-cyclopent-3-enyl)-but-2-enyl ester was prepared from 3-(2-hydroxy-5-methyl-phenyl)-acrylic acid ethyl ester, 2-ethyl-4(2,2,3-

trimethyl cyclopentyl-3-en-1-yl)-but-2-en-1-ol and tetraisopropyl-o-titanate.

5 (E)-3-(2-Hydroxy-5-methyl-phenyl)-acrylic acid 3-methyl-5-phenyl-pentyl ester

According to the same procedure, 3-(2-hydroxy-5-methyl-phenyl)-acrylic acid 3-methyl-5-phenyl-pentyl ester was prepared from 3-(2-hydroxy-5-methyl-phenyl)-acrylic acid ethyl ester, 3-methyl-5-phenyl-pentanol and tetraisopropyl-o-titanate.

15 (E)-3-(4-Diethylamino-2-hydroxy-phenyl)-but-2-enoic acid dec-9-enyl ester

According to the same procedure, 3-(4-diethylamino-2-hydroxy-phenyl)-but-2-enoic acid dec-9-enyl ester was prepared from 3-(4-diethylamino-2-hydroxy-phenyl)-but-2-enoic acid ethyl ester, dec-9-en-1-ol and tetraisopropyl-o-titanate.

20 (E)-3-(4-Diethylamino-2-hydroxy-phenyl)-but-2-enoic acid 3-methyl-5-phenyl-pentyl ester

According to the same procedure, 3-(4-diethylamino-2-hydroxy-phenyl)-but-2-enoic acid 3-methyl-5-phenyl-pentyl ester was prepared from 3-(4-diethylamino-2-hydroxy-phenyl)-but-2-enoic acid ethyl ester, 3-methyl-5-phenyl-pentanol and tetraisopropyl-o-titanate.

30 Example 9

3-[2-(tert-Butyl-dimethyl-silyloxy)-phenyl]-acrylic acid
tert-butyl-dimethyl-silyl ester

To a solution of 5.0 g of (E)-3-(2-hydroxy-phenyl)-acrylic acid and 4.6 g of imidazole in 100 ml of DMF was added at room temperature a solution of 10.1 g of TBDMS-Cl. The reaction mixture was stirred overnight, poured onto 200 ml of 5 cold water and extracted 3x with 150 ml of MTBE. The combined organic phases were dried ($MgSO_4$) and evaporated to dryness. The resulting oil was dried at 0.08 Torr/120-170°C to remove excess of TBDMS-Cl to yield 11.4 g of 3-[2-(tert-butyl-dimethyl-silanyloxy)-phenyl]-acrylic acid tert-butyl-dimethyl-10 silyl ester as a yellowish oil.

NMR ($CDCl_3$) δ 7.96 (d, 1H), 7.52 (dd, 1H), 7.23 (m, 1H), 6.94 (m, 1H), 6.82 (dd, 1H), 6.37 (d, 1H), 1.00 (s, 9H), 0.97 (s, 9H), 0.32 (s, 6H), 0.22 (s, 6H)

15

Example 10

3-[2-(tert-Butyl-dimethyl-silanyloxy)-phenyl]-acryloyl
chloride

20 To a 0°C cold solution of 6.3 g of 3-[2-(tert-Butyl-dimethyl-silanyloxy)-phenyl]-acrylic acid tert-butyl-dimethyl-silyl ester and 8 drops of DMF in 15 ml of CH_2Cl_2 was added dropwise 2.0 ml of oxalyl chloride. After complete addition the reaction mixture was allowed to warm to room temperature 25 and stirring was continued for 60 h. The reaction mixture was filtered, evaporated to dryness and taken up in MTBE. The solution was cooled in the refrigerator overnight, filtered and evaporated to yield 4.4 g of 3-[2-(tert-Butyl-dimethyl-silanyloxy)-phenyl]-acryloyl chloride.

30

NMR ($CDCl_3$) δ 8.28 (d, 1H), 7.55 (dd, 1H), 7.35 (m, 1H), 7.01 (m, 1H), 6.87 (dd, 1H), 6.63 (d, 1H), 1.04 (s, 9H), 0.26 (s, 6H)

Example 11

3-(2-Hydroxy-phenyl)-acrylic acid 3-(3-isopropyl-phenyl)-but-1-enyl ester

5

To a suspension of 0.72 g of NaH (55% in oil), previously washed with hexane, in 22.5 ml of THF was added 1.67 ml of tert-butanol over a period of 12 min. The reaction mixture was stirred at room temperature for 1.25 h, was then cooled to -15°C and 2.85 g of 3-(3-isopropyl-phenyl)-butanal (Florhydral) was slowly added. Stirring was continued for 45 min. and then this cold enolate solution was added via Teflon canula to a -5°C cold solution of 4.44 g of 3-[2-(tert-butyl-dimethyl-silyloxy)-phenyl]-acryloyl chloride in 7.5 ml of THF. After complete addition stirring at -5°C was continued for 1 h, the reaction mixture was quenched with 80 ml of water / 40 ml of brine and was extracted with MTBE. The combined organic phases were washed with water/brine 2:1, dried ($MgSO_4$) and evaporated to dryness to yield 6.65 g of crude 3-[2-(tert-Butyl-dimethyl-silyloxy)-phenyl]-acrylic acid 3-(3-isopropyl-phenyl)-but-1-enyl ester as a yellowish, viscous oil.

To a 0°C cold solution of 6.2 g of this oil in 70 ml of THF was added slowly 13.7 ml of a 1M TBAF/THF solution. The reaction mixture was stirred at 0°C for 1.5 h, poured onto 200 ml of H_2O and extracted 3x with 150 ml of MTBE. The combined organic phases were washed with brine, dried ($MgSO_4$) and evaporated to dryness. The resulting oil was purified by chromatography to yield 2.64 g of 3-(2-Hydroxy-phenyl)-acrylic acid 3-(3-isopropyl-phenyl)-but-1-enyl ester in form of a yellowish oil.

NMR ($CDCl_3$) δ 8.12 and 8.01 (d, 1H), 7.50-6.80 (m, 10H), 6.66 and 6.56 (d, 1H), 5.75 and 5.13 (m, 1H), 4.13

(m, CH), 3.53 (m, CH), 2.89 (m, CH), 1.42 (d, 3H), 1.25 (d, 6H)

Example 12

5 (E)-3-[2-(tert-Butyl-dimethyl-silanoxy)-phenyl]-acrylic acid

To a solution of 8.2 g of (E)-3-(2-hydroxy-phenyl)-acrylic acid and 15.2 g of TBDMS-Cl in 10 ml of DMF was added 11.9 g of imidazole and the mixture was stirred at room 10 temperature for 24 h. After aqueous work-up, the bis-silylated product was saponified with 2.1 g of LiOH in THF-H₂O for 0.5 h at 0°C. The mixture was concentrated *in vacuo* and TBDMS-OH removed by extraction with hexane. The aqueous layer was acidified to pH 4 with KHSO₄ and extracted with EtOAc. The 15 organic phases were washed with brine, dried (Na₂SO₄) and evaporated to dryness. The semi-solid residue was suspended in hexane, filtered and the filtrate evaporated to dryness to yield 7.5 g of a pale yellow oil, which solidified upon standing.

20

NMR (CDCl₃) δ 8.08 (d, 1H), 7.46 (dd, 1H), 7.18 (dt, 1H), 6.87 (t, 1H), 6.75 (dd, 1H), 6.31 (d, 1H), 0.93 (s, 9H), 0.13 (s, 6H)

25

Example 13

(E)-3-(2-Hydroxy-phenyl)-acrylic acid 1-ethoxy-3-(3-isopropyl-phenyl)-butyl ester

A solution of 6.5 g of (E)-3-[2-(tert-Butyl-dimethyl-silanoxy)-phenyl]-acrylic acid and 5.5 g of 1-(3-ethoxy-1-methyl-allyl)-3-isopropyl-benzene, accessible from 3-(3-isopropyl-phenyl)-butyraldehyde via a two step procedure according to P.D. Bartlett and A.A. Frimer, *Heterocycles* 11, 419-435, (1978), in 25 ml of toluene was heated to reflux and

stirred for 16 h. After concentration, the reaction mixture was purified by flash chromatography to yield 7.5 g of the intermediate (E)-3-[2-(*tert*-Butyl-dimethyl-silanoxy)-phenyl]-acrylic acid 1-ethoxy-3-(3-isopropyl-phenyl)-butyl ester as a 5 pale yellow oil. This was dissolved in 50 ml THF and treated at 0°C with 4.7 g of TBAF•3H₂O. After 20 min., the mixture was concentrated and purification by flash chromatography yielded 2.2 g of the title ester (a mixture of diastereoisomers) as a pale yellow oil.

10

NMR (CDCl₃) δ 8.12 and 8.08 (d, 1H), 7.47 (dt, 1H), 7.35 (br s, 1H), 7.31-7.18 (m, 2H), 7.12-6.98 (m, 3H), 6.92 (dt, 1H), 6.83 (dd, 1H), 6.68 and 6.61 (d, 1H), 5.96 and 5.77 (dd, 1H), 3.82-3.65 (m, 1H), 3.63-3.40 (m, 1H), 3.05-2.77 (m, 2H), 2.22-1.95 (m, 2H) and 1.38-1.12 (m, 12H)

Example 14

(E)-3-(2-Hydroxy-phenyl)-acrylic acid 3-(4-*tert*-butyl-phenyl)-1-ethoxy-propyl ester

A solution of 3.1 g of (E)-3-[2-(*tert*-Butyl-dimethyl-silanoxy)-phenyl]-acrylic acid, 3.0 g of 1-*tert*-butyl-4-(3-ethoxy-2-methyl-allyl)-benzene, accessible from 3-(4-*tert*-butyl-phenyl)-2-methyl-propionaldehyde via a two step 25 procedure according to P.D. Bartlett and A.A. Frimer, Heterocycles 11, 419-435, (1978), and 20 mg of *p*-TSA in 25 ml of toluene was stirred at 0 °C for 4 h and at room temperature for 14 h. The reaction mixture was partitioned between 30 saturated sodium carbonate and hexane and the organic layer was washed with brine, dried over (Na₂SO₄) and evaporated to dryness to yield 5.3 g of the crude intermediate (E)-3-[2-(*tert*-Butyl-dimethyl-silanoxy)-phenyl]-acrylic acid 3-(4-*tert*-butyl-phenyl)-1-ethoxy-propyl ester as a pale yellow oil. This

was dissolved in 20 ml of THF and treated at 0 °C with 3.2 g of solid TBAF•3H₂O. After 30 min., the mixture was concentrated and purification by repeated flash chromatography yielded 1.2 g of the title ester (a mixture of 5 diastereoisomers) as a pale yellow oil.

NMR (CDCl₃) d 8.17 and 8.13 (d, 1H), 7.50 (ddd, 1H), 7.36-10 7.18 (m, 4H), 7.11 (br dd, 2H), 6.98-6.83 (m, 2H), 6.73 and 6.69 (d, 1H), 5.94 (t, 1H), 3.92-3.70 (m, 1H), 3.71-3.54 (m, 1H), 3.05-2.81 (m, 1H), 2.52-2.34 (m, 1H), 2.30-2.12 (m, 1H) and 1.31 (s, 9H), 1.33-1.20 (m, 3H), 0.97 and 0.90 (t, 3H)

15 **Example 15**

Test cloth was washed with detergent to which one or more of the precursors of Examples 1-14 had been added. The cloth was then line dried. The cloth dried in sunlight had a 20 distinct fragrance note, as determined by a trained panel. In contrast, the cloth dried without sunlight was olfactively neutral.

25 **Example 16**

Test cloth was washed with a detergent and then a fabric softener, containing one ore more of the precursors of Examples 1-14, was added to the rinse cycle. The cloth was then line dried. The cloth dried in sunlight had a distinct 30 fragrance note, as determined by a trained panel. In contrast, the cloth dried without sunlight was olfactively neutral.

Example 17

A 1% solution of one or more of the products of Examples 1-14 in ethanol was applied to cigarette papers to produce levels of 5-50'000 ppm of each flavorant. The paper was incorporated in cigarettes and, upon burning, released a
5 fragrant odor.

Example 18

10

A broad spectrum (UV-A and UV-B) oil/water sunscreen lotion was prepared with 0,5%.

Part A

	<u>Recipe: %</u>	<u>Compound</u>	<u>Chemical Name</u>
15	2 %	PARSOL MCX	Octyl methoxycinnamate
	3 %	PARSOL 1789	4-4-Butyl-4'methoxy-dibenzoyl methane
20	12 %	Cétiol LC	Coco-caprylate/caprate
	4 %	Dermol 185	Isostearyl neopentanoate
	0,25 %	Diethyl- eneglycol monostearate	PEG-2-stearate
	1 %	Cetylalcohol	Cetylalcohol
25	0,25 %	MPOB/PPOB	Methyl-propylparabene
	0,1 %	EDTA BD	EDTA-sodium salt
	1 %	Amphisol DEA (Giv.)	Diethanolamine cetyl-phosphate

30

Part B

20 %	Permulene TR-1 (+%)	Acrylate C10-C30 Alkyl-acrylate
50,1 %	water deion	water deion

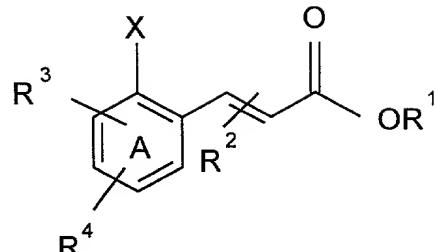
5 %	Propyleneglycol 1,2-Propanediol
0,8 %	KOH (10%) Potassium hydroxide

Part A was heated in a reactor to 85°C. Part B was
5 slowly added within 10 min., followed by addition of KOH and
0.5% of the product of Example 5. The emulsion was then cooled
and degassed.

The invention being thus described, it will be obvious
10 that the same may be varied in many ways. Such variations are
not to be regarded as a departure from the spirit and scope of
the invention and all such modifications are intended to be
included within the scope of the following claims.

What Is Claimed Is:

1. A compound of Formula I



5

wherein

10 A is selected from benzene and naphthalene rings;

15 R¹ is a saturated, unsaturated, straight, branched, alicyclic or an aromatic C₁₀-C₃₀ hydrocarbon residue which can contain heteroatoms and can be substituted by an ionic substituent;

20 R² in 2- or 3-position is a hydrogen, a straight or branched C₁-C₆ residue, an optionally substituted aromatic or an optionally substituted heterocyclic residue;

25 R³ and R⁴ are a hydrogen, a straight or branched C₁-C₆ alkyl, a C₁-C₆ alkoxy residue, a substituted or condensed heterocyclic residue, -OH, -NO₂, -NH₂, -N(C₁-C₆ alkyl)₂, -N(hydroxyalkyl)₂, -NHCO₂CH₃ or -NH(heterocycle),

wherein R², R³ and R⁴ are the same or different;

X is an -OH or NHR⁶, wherein R⁶ is a hydrogen, a saturated or unsaturated, straight or branched C₁-C₂₀ hydrocarbon, or

an optionally substituted aromatic or heterocyclic residue;

and the acrylic double bond is of the E configuration.

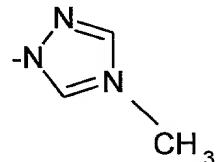
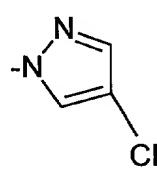
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2. A compound according to claim 1 wherein R¹ is a saturated, unsaturated, straight or branched C₁₀-C₃₀ hydrocarbon residue comprising one or more O atoms, N atoms, C(O) groups, alkoxy groups and mixtures thereof.

10

3. A compound according to claim 1 wherein R¹ is a saturated, unsaturated, straight or branched C₁₀-C₃₀ hydrocarbon residue substituted by an ionic substituent of Formula NR⁵₃⁺, wherein R⁵ is a residue of a fatty acid or an alkyl group with 1 to 30 carbon atoms.

4. A compound according to claim 1 wherein R² is a heterocyclic residue of Formula



20

5. A compound according to claim 1 wherein at least one of R³ and R⁴ is a five membered heterocyclic residue comprising N atoms and/or O atoms.

25

6. A compound according to claim 1 wherein at least one of R³ and/or R⁴ is a hydrogen, -N(C₁-C₆ alkyl)₂, -NH₂, or a five membered heterocyclic residue, substituted by C₁-C₆ aliphatic and/or aromatic substituents.

30

7. A compound according to claim 1 wherein R^2 is hydrogen or methyl.

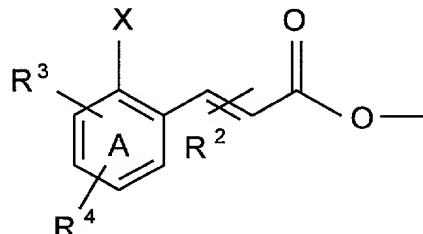
8. A compound according to claim 1 wherein R^1 is a residue of
5 an olfactory alcohol of Formula R^1OH .

9. A compound according to claim 1 wherein R^1 is a residue of the enol form of an olfactory aldehyde of Formula R^1HO .

10 10. A compound according to claim 1 wherein R^1 is a residue of the enol form of an olfactory ketone of Formula R^1O .

11. A compound according to claim 1 wherein R^1 is a substituted alkyl, an alkenyl or an arylalkyl residue
15 carrying a 1-alkoxy, 1-aryloxy or 1-arylalkoxy residue.

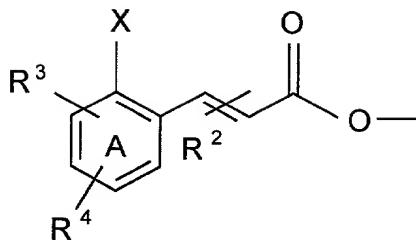
12. A compound according to claim 1 wherein a residue of Formula Ia:



20

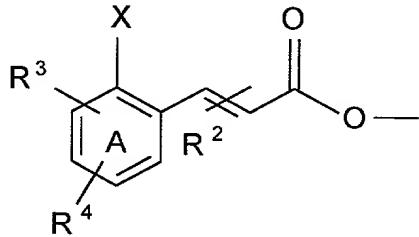
is a precursor for a fragrant coumarin.

13. A compound according to claim 1 wherein a residue of
25 Formula Ia



is a precursor for a fluorescent whitening coumarin.

5 14. A compound according to claim 12 wherein R¹ is a residue
of an olfactory alcohol, an aldehyde or ketone and the
residue of Formula Ia

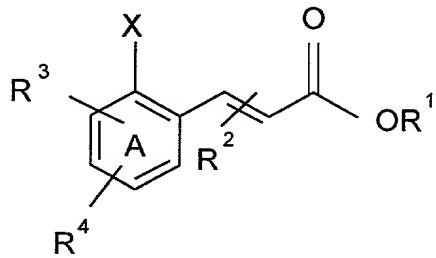


10

is a precursor for a fragrant coumarin.

15 15. A method for preparing compositions which provide upon
activation organoleptic, antimicrobial or fluorescent
whitening properties comprising incorporating into one of
these compositions a compound of Formula I:

(I)



20 wherein

A is selected from benzene and naphthalene rings;

5 R¹ is a saturated, unsaturated, straight, branched, alicyclic or an aromatic C₁₀-C₃₀ hydrocarbon residue which can contain heteroatoms and can be substituted by an ionic substituent;

10 R² in the 2- or 3-position is a hydrogen, a straight or branched C₁-C₆ residue, an optionally substituted aromatic or an optionally substituted heterocyclic residue;

15 R³ and R⁴ are a hydrogen, a straight or branched C₁-C₆ alkyl, a C₁-C₆ alkoxy residue, a substituted or condensed heterocyclic residue, -OH, -NO₂, -NH₂, -N(C₁-C₆ alkyl)₂, -N(hydroxyalkyl)₂, -NHCO₂CH₃ or -NH(heterocycle);

wherein R², R³ and R⁴ are the same or different;

20 X is an -OH or NHR⁶, wherein R⁶ is a hydrogen, a saturated or unsaturated, straight or branched C₁-C₂₀ hydrocarbon, or an optionally substituted aromatic or heterocyclic residue; and

25 the acrylic double bond is of the E configuration.

16. A method according to claim 15 wherein the precursors are incorporated into laundry products.

30 17. A method according to claim 15 wherein the precursors are incorporated into tobacco products.

18. A method according to claim 15 wherein the precursors are incorporated into cosmetics and toiletries.

ABSTRACT

The acrylic acid esters of Formula I are useful for the delivery of organoleptic compounds, especially for flavors, 5 fragrances, masking agents and antimicrobial compounds. They can also deliver fluorescent whitening agents.

**DECLARATION AND POWER OF ATTORNEY
FOR UTILITY PATENT APPLICATION - JOINT**

As the below named inventors, we hereby declare that :

Our residences, post office addresses, and citizenships are as stated below next to our names.

We believe that we are the original, first, and joint inventors of the subject matter which is claimed and for which a patent is sought on the invention entitled **Aryl-acrylic acid esters**, the specification of which is attached hereto.

We hereby state that we reviewed and understand the contents of the above-identified specification, including the claims, as amended by any amendment referred to above.

We acknowledge the duty to disclose information that is material to the examination of this application in accordance with 37 Code of Federal Regulations § 1.56.

We hereby claim foreign priority benefits under 35 United States Code §§ 119 and 365 of any foreign application(s) for the patent or inventor's certificate listed below and have also identified below any foreign application for patent or inventor's certificate having a filing date before that of the application on which priority is claimed :

Prior Foreign Application :

European Patent Application No. 98810114.3, filed February 13, 1998

We hereby appoint the following attorneys to prosecute this application and to transact all business in the Patent and Trademark Office connected herewith :

Maurice B. Stiefel, Reg. No. 18, 479; Lawrence G. Kurland, Reg. No. 24,895; Stephen P. Gilbert, Reg. No. 27,893; Mark E. Waddell, Reg. No. 31,803, David A. Roodman, Reg. No. 35,663; Stephen M. Haracz, Reg. No. 33,397; N. Whitney Wilson, Reg. No. 38,661; Daniel A. Crowe, Reg. No. 39,644; Timothy E. Tracy, Reg. No. 39,401; Warren K. MacRae, Reg. No. 37,876; Leo G. Lenna, Reg. No. P-42,796; Kathleen Gribschaw, Reg. No. P-41,806, Todd S. Sharinn, Reg. No. P-42,144; Robert J. Lipka, Reg. No. P-42,807; and Christopher J. Hayes, Reg. No. P-41,822.

Address all telephone calls to Mark E. Waddell, Esq., or alternatively to Eileen M. Ebel, Esq., at telephone number (212) 692-1800.

Address all correspondence to :

Mark E. Waddell, Esq.
BRYAN CAVE LLP
245 Park Avenue
New York, New York 10167-0034

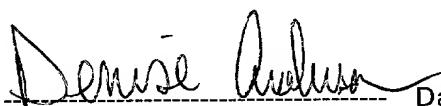
We hereby declare that all statements made herein of our own knowledge are true and that all statements made on information and belief are believed to be true; and further these statements were made with the knowledge that willful false statements and the likes so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full Name of First Inventor : ANDERSON Denise, Dr.

Residence : Zürich, Switzerland

Citizenship : USA

Post Office Address : Hottingerstrasse 5
CH-8032 Zürich
Switzerland

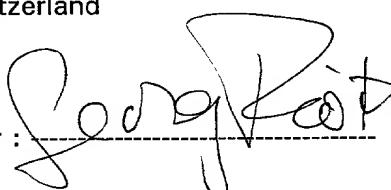
Signature of First Inventor :  Date : 01/18/1993

Full Name of Second Inventor : FRATER Georg, Dr.

Residence : Winterthur, Switzerland

Citizenship : Swiss

Post Office Address : Turmstrasse 61
CH-8400 Winterthur
Switzerland

Signature of Second Inventor :  Date : 01-18-1993